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and Con, trol, Blanche Lane, South Mimms, Potters Bar,  
Hertfordshire EN6 2QG (GB).

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(74) Agent: I.P. 21 LIMITED; Norwich Research Park, Col-  
ney, Norwich NR4 7UT (GB).

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(71) Applicant (*for all designated States except US*): NA-  
TIONAL BIOLOGICAL STANDARDS BOARD  
[GB/GB]; National Institute for Biological Standards and  
Con, trol, Blanche Lane, South Mimms, Potters Bar,  
Hertfordshire EN6 2QG (GB).

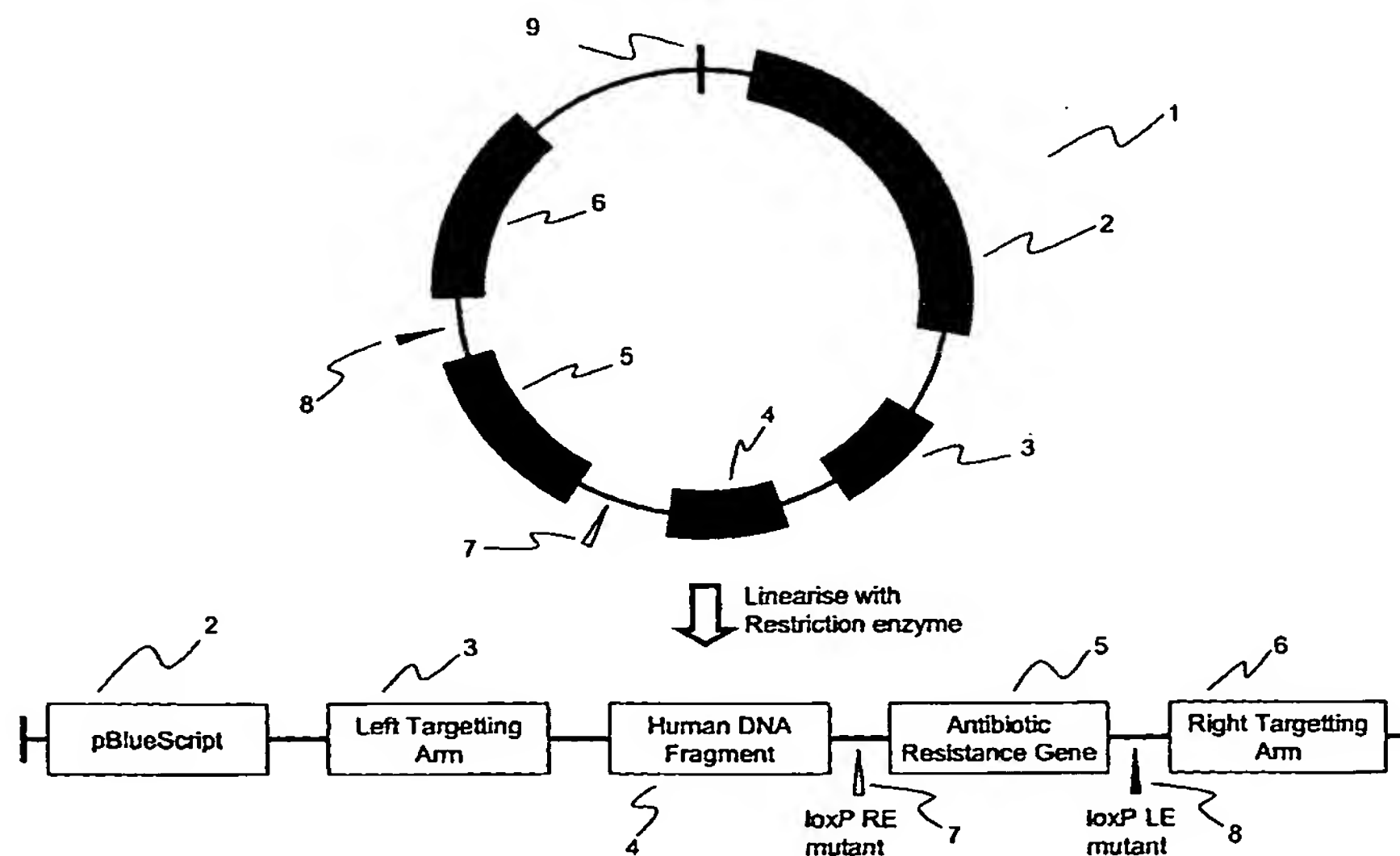
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(72) Inventor; and

(75) Inventor/Applicant (*for US only*): HAWKINS, John,  
Ross [GB/GB]; National Institute for Biological Standards

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(54) Title: GENETIC REFERENCE MATERIALS



(57) Abstract: The invention provides a genetic reference standard comprising at least one human genetic reference sequence (comprising a human DNA sequence containing at least one genetic variant whose presence in the DNA of a human subject is indicative of a pathological condition, a predisposition to a pathological condition, or a predisposition to an adverse reaction to external stimuli, or is indicative of a patient's likely response to a therapeutic intervention, i.e. a variant used in pharmacogenomic analysis) cloned into a non-mammalian animal cell line. There are also provided such reference standards where the human DNA is targeted to specific location in the host genome, using homologous recombination. The invention further provides a method of detecting a genetic variant using such reference standards.

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